

## TENT COOPERATION TRE Y

PCT

## NOTIFICATION OF ELECTION

(PCT Rule 61.2)

From the INTERNATIONAL BUREAU

To:

Assistant Commissioner for Patents  
United States Patent and Trademark  
Office  
Box PCT  
Washington, D.C.20231  
ETATS-UNIS D'AMERIQUE

in its capacity as elected Office

<b>Date of mailing (day/month/year)</b> 13 June 2000 (13.06.00)	
<b>International application No.</b> PCT/US99/17167	<b>Applicant's or agent's file reference</b> PF-0568 PCT
<b>International filing date (day/month/year)</b> 30 July 1999 (30.07.99)	<b>Priority date (day/month/year)</b> 31 July 1998 (31.07.98)
<b>Applicant</b> BANDMAN, Olga et al	

1. The designated Office is hereby notified of its election made:

☒ in the demand filed with the International Preliminary Examining Authority on:

02 February 2000 (02.02.00)

☐ in a notice effecting later election filed with the International Bureau on:
2. The election ☒ was
☐ was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

<b>The International Bureau of WIPO</b> 34, chemin des Colombettes 1211 Geneva 20, Switzerland Facsimile No.: (41-22) 740.14.35	<b>Authorized officer</b>  Kiwa Mpay Telephone No.: (41-22) 338.83.38
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# PATENT COOPERATION TREATY

From the  
INTERNATIONAL PRELIMINARY EXAMINING AUTHORITY

To: LUCY J. BILLINGS  
INCYTE PHARMACEUTICALS, INC.  
3174 PORTER DRIVE  
PALA ALTO, CA 94304

## PCT

### NOTIFICATION OF TRANSMITTAL OF INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Rule 71.1)

Date of Mailing  
(day/month/year)

22 JAN 2001

Applicant's or agent's file reference

PF-0568 PCT

IMPORTANT NOTIFICATION

International application No.

PCT/US99/17167

International filing date (day/month/year)

30 JULY 1999

Priority Date (day/month/year)

31 JULY 1998

Applicant

INCYTE PHARMACEUTICALS, INC.

1. The applicant is hereby notified that this International Preliminary Examining Authority transmits herewith the international preliminary examination report and its annexes, if any, established on the international application.
2. A copy of the report and its annexes, if any, is being transmitted to the International Bureau for communication to all the elected Offices.
3. Where required by any of the elected Offices, the International Bureau will prepare an English translation of the report (but not of any annexes) and will transmit such translation to those Offices.
4. **REMINDER**

The applicant must enter the national phase before each elected Office by performing certain acts (filing translations and paying national fees) within 30 months from the priority date (or later in some Offices)(Article 39(1))(see also the reminder sent by the International Bureau with Form PCT/IB/301).

Where a translation of the international application must be furnished to an elected Office, that translation must contain a translation of any annexes to the international preliminary examination report. It is the applicant's responsibility to prepare and furnish such translation directly to each elected Office concerned.

For further details on the applicable time limits and requirements of the elected Offices, see Volume II of the PCT Applicant's Guide.

Name and mailing address of the IPEA/US

Commissioner of Patents and Trademarks  
Box PCT  
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

KAREN COCHRANE CARLSON, PH.D.

Telephone No. (703) 308-0116

**DELLA MAE COLLINS**  
**PARALEGAL SPECIALIST**  
**TECHNOLOGY CENTER 1600**

# PATENT COOPERATION TREATY

## PCT

### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference PF-0568 PCT	<b>FOR FURTHER ACTION</b> See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)	
International application No. PCT/US99/17167	International filing date (day/month/year) 30 JULY 1999	Priority date (day/month/year) 31 JULY 1998
International Patent Classification (IPC) or national classification and IPC Please See Supplemental Sheet.		
Applicant INCYTE PHARMACEUTICALS, INC.		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.
2. This REPORT consists of a total of 4 sheets.  
☐ This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority. (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

These annexes consist of a total of 0 sheets.

3. This report contains indications relating to the following items:

- I ☒ Basis of the report
- II ☐ Priority
- III ☐ Non-establishment of report with regard to novelty, inventive step or industrial applicability
- IV ☐ Lack of unity of invention
- V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI ☐ Certain documents cited
- VII ☐ Certain defects in the international application
- VIII ☐ Certain observations on the international application

Date of submission of the demand  02 FEBRUARY 2000	Date of completion of this report  06 DECEMBER 2000
Name and mailing address of the IPEA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231  Facsimile No. (703) 305-3230	Authorized officer  KAREN COCHRANE Telephone No. (703) 308-0196

## INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/US99/17167

## I. Basis of the report

## 1. With regard to the elements of the international application:\*

☒ the international application as originally filed☒ the description:

pages 1-58 , as originally filed  
pages NONE , filed with the demand  
pages NONE , filed with the letter of \_\_\_\_\_

☒ the claims:

pages 1-20 , as originally filed  
pages NONE , as amended (together with any statement) under Article 19  
pages NONE , filed with the demand  
pages NONE , filed with the letter of \_\_\_\_\_

☒ the drawings:

pages 1 , as originally filed  
pages NONE , filed with the demand  
pages NONE , filed with the letter of \_\_\_\_\_

☒ the sequence listing part of the description:

pages 1-21 , as originally filed  
pages NONE , filed with the demand  
pages NONE , filed with the letter of \_\_\_\_\_

2. With regard to the language, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.  
These elements were available or furnished to this Authority in the following language \_\_\_\_\_ which is:

- ☐ the language of a translation furnished for the purposes of international search (under Rule 23.1(b)).  
☐ the language of publication of the international application (under Rule 48.3(b)).  
☐ the language of the translation furnished for the purposes of international preliminary examination (under Rules 55.2 and/or 55.3).

## 3. With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:

- ☐ contained in the international application in printed form.  
☐ filed together with the international application in computer readable form.  
☐ furnished subsequently to this Authority in written form.  
☐ furnished subsequently to this Authority in computer readable form.  
☐ The statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.  
☐ The statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished.

4. ☒ The amendments have resulted in the cancellation of:

- ☒ the description, pages NONE  
☒ the claims, Nos. NONE  
☒ the drawings, sheets/fig NONE

5. ☐ This report has been drawn as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).\*\*

\* Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17).

\*\*Any replacement sheet containing such amendments must be referred to under item 1 and annexed to this report.

# INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/US99/17167

## V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

### 1. statement

Novelty (N)	Claims	<u>13, 14, 16-20</u>	YES
	Claims	<u>1-12, 15</u>	NO
Inventive Step (IS)	Claims	<u>13, 14, 16-20</u>	YES
	Claims	<u>1-12, 15</u>	NO
Industrial Applicability (IA)	Claims	<u>1-20</u>	YES
	Claims	<u>NONE</u>	NO

### 2. citations and explanations (Rule 70.7)

Claims 1-12 and 15 lack novelty under PCT Article 33(2) as being anticipated by Ryseck et al. Ryseck et al teach V58 that is 92% identical to SEQ ID NO:1 (Claims 1, 2). Nucleic acid encoding V58 is shown in Fig. 1 (Claims 6, 9-11, 15). Northern blot analysis using the polynucleotide encoding V58 is shown (Claim 7) in which AMV reverse transcriptase was used to amplify the polynucleotide prior to Dot blot analysis (Claim 8). The polynucleotides were transformed into M13-derived vectors (Claim 12) or pUC19 plasmids.

Claims 13, 14, and 16-20 meet the criteria set out in PCT Article 33(2)-(4), because the prior art does not teach or fairly suggest a host cell comprising the vector having polynucleotide encoding SEQ ID NO:1, the recombinant production of the polypeptide, antibody, agonist, or antagonist to the polypeptide, or a method of treatment.

A response to this written opinion was received 15 Nov 2000. Applicants made no direct argument against the opinion, stating only that they elect to reserve the right to address objections in national stage applications.

----- NEW CITATIONS -----  
NONE

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No.

PCT/US99/17167

**Supplemental Box**

(To be used when the space in any of the preceding boxes is not sufficient)

Continuation of: Boxes I - VIII

Sheet 10

**CLASSIFICATION:**

The International Patent Classification (IPC) and/or the National classification are as listed below:

IPC(7): C07K14/00; C07H 17/00 C12P 14/00 and US Cl.: 435/ 69.1. 252.3, 320.1, 325; 536/23.1; 530/350

# PATENT COOPERATION TREATY

## PCT

### INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

REC'D 25 JAN 2001

WIPO

PCT

15

Applicant's or agent's file reference PF-0568 PCT	<b>FOR FURTHER ACTION</b> See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)	
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Novelty (N)

Claims 13, 14, 16-20

YES

Claims 1-12, 15

NO

Inventive Step (IS)

Claims 13, 14, 16-20

YES

Claims 1-12, 15

NO

Industrial Applicability (IA)

Claims 1-20

YES

Claims NONE

NO

### 2. citations and explanations (Rule 70.7)

Claims 1-12 and 15 lack novelty under PCT Article 33(2) as being anticipated by Ryseck et al. Ryseck et al teach V58 that is 92% identical to SEQ ID NO:1 (Claims 1, 2). Nucleic acid encoding V58 is shown in Fig. 1 (Claims 6, 9-11, 15). Northern blot analysis using the polynucleotide encoding V58 is shown (Claim 7) in which AMV reverse transcriptase was used to amplify the polynucleotide prior to Dot blot analysis (Claim 8). The polynucleotides were transformed into M13-derived vectors (Claim 12) or pUC19 plasmids.

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